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In the Claims:

For the Examiner's convenience, Applicants present all pending claims with status

indicators in compliance with the practice guidelines for making amendments under 37

C.F.R. §1.121(c)(1).

Please cancel claims 1-29 and 32. Additionally, please amend claims 30, 33-35 and add

new claims 36-37 as follows:

Claims 1-29 CANCEL

30. (CURRENTLY AMENDED) A method of detecting a disease-state in a subject,

wherein the disease-state is associated with expression of an RG1 polypeptide

having the amino acid sequence of SEQ ID NO: 2, and wherein the method

comprises:

(a) administering to the subject an immunoconjugate-of Claim 22, wherein the

immunoconjugate comprises an antibody or a fragment or a variant thereof

that specifically binds to an epitope present in the RG1 polypeptide having

the amino acid sequence of SEQ ID NO: 2 and wherein the antibody or the

fragment or the variant thereof is conjugated to a molecule which is a

detectable marker;

(b) detecting the binding of the immunoconjugate within the subject.; and

(c) determining if the level of binding of the immunoconjugate in the subject

is increased as compared with the level of binding detected in disease-free

control subjects, an increased level being indicative of a disease state.

31. (PREVIOUSLY PRESENTED) The method of Claim 30, wherein the method of

detection is immunoscintigrapy.

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- 32. (CANCEL)
- 33. (CURRENTLY AMENDED) The method of Claim 32 30, wherein the method of detection is positron emitting tomography.
- 34. (CURRENTLY AMENDED) The method of Claim 33-30, wherein the detectable marker of the immunoconjugate is selected from a group consisting of ⁴³Sc, ⁴⁴Sc, ⁵²Fe, ⁵⁵Co, ⁶⁸Ga, ⁶⁴Cu, ⁸⁶Y, or ^{94m}Tc, ¹¹¹In, or ⁹⁰Y.
- 35. (CURRENTLY AMENDED) The method of Claim 34-30, wherein the disease-state is prostate cancer.
- 36. (NEW) The method of claim 30, wherein the detectable marker is a radioisotope, a fluorescent compound, a bioluminescent compound, chemiluminescent compound, a metal chelator or an enzyme.
- 37. (NEW) The method of claim 30, wherein the antibody is selected from a group consisting of a polyclonal antibody, a monoclonal antibody, a chimeric antibody, a humanized antibody and a fully-human antibody.